

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of claims:

1. - 5. (Canceled)

6. (Currently amended) A pharmaceutical composition comprising a fixed combination of active compounds consisting of the active compound ciclesonide or a solvate, an epimer, physiologically functional derivative or a solvate of a physiologically functional derivative thereof and the active compound R,R-formoterol or a ~~hydrate, solvate, salt, or~~ hydrate of a salt ~~or solvate of a salt~~ thereof, ~~in fixed combination and in an administration form suitable for inhalative administration by means of a powder inhaler,~~

wherein the active compound ciclesonide or a solvate, an epimer, physiologically functional derivative or solvate of a physiologically functional derivative thereof and the active compound R,R-formoterol or a ~~hydrate, solvate, salt, or~~ hydrate of a salt ~~or solvate of a salt~~ thereof, are present ready mixed in a fixed combination.

7. (Previously presented) The pharmaceutical composition according to Claim 6, wherein the active compound ciclesonide is present as its R epimer in an amount greater than 95%.

8. (Currently amended) The pharmaceutical composition according to Claim 6, wherein the active compound ciclesonide is present as its R epimer in an amount greater than 95%, and the active compound R,R-formoterol is present as a ~~hydrate~~, salt or hydrate of a salt thereof.

9. (Previously presented) The pharmaceutical composition according to Claim 8, wherein the active compound R,R-formoterol is present as a salt with an acid selected from the group consisting of hydrochloric acid, hydrobromic acid, phosphoric acid, nitric acid, sulphuric acid, acetic acid, citric acid, D-gluconic acid, benzoic acid, 2-(4-hydroxybenzoyl) benzoic acid, butyric acid, sulphosalicylic acid, maleic acid, lauric acid, malic acid, fumaric acid, succinic acid, oxalic acid, tartaric acid, embonic acid, stearic acid, toluenesulphonic, methanesulphonic acid and 1-hydroxy-2-naphthoic acid.

10. (Previously presented) The pharmaceutical composition according to claim 9, wherein the acid is fumaric acid or tartaric acid.

11. (Currently amended) A method of treating an airway disease in a patient comprising administering to a patient in need thereof a pharmaceutical composition comprising a fixed combination of active compounds consisting of a therapeutically effective amount of the active compound ciclesonide or a solvate, an epimer, physiologically functional derivative or solvate of a physiologically functional derivative thereof and the active compound R,R-formoterol or a salt or

hydrate of a salt thereof, in fixed combination and in an administration form suitable for inhalative administration by means of a powder inhaler with the active compound R,R-formoterol or a hydrate, solvate, salt, hydrate of a salt or solvate of a salt thereof, wherein the active compound ciclesonide or a solvate, an epimer, physiologically functional derivative or solvate of a physiologically functional derivative thereof and the active compound R,R-formoterol or a hydrate, solvate, salt, or hydrate of a salt or solvate of a salt thereof, are present ready mixed in a fixed combination.

12. (Canceled)

13. (Previously presented) The method according to Claim 11, wherein the active compound ciclesonide is present as its R epimer in an amount greater than 95%.

14. (Previously presented) A method of treating an airway disease in a patient comprising administering to a patient in need thereof a therapeutically effective amount of the pharmaceutical composition according to claim 6 by means of a dry powder inhaler.

15. (Previously presented) The method according to claim 14, wherein the airway disease is selected from the group consisting of bronchitis, obstructive bronchitis, COPD (chronic obstructive pulmonary disease), spastic bronchitis, allergic bronchitis, allergic asthma and bronchial asthma.

16. (Currently amended) The method according to claim 14, wherein ciclesonide or a solvate, an epimer, physiologically functional derivative or solvate of a physiologically functional derivative thereof is administered in a dose of 0.05 to 1 mg per day and R,R-formoterol or a hydrate, solvate, salt, or hydrate of a salt or solvate of a salt thereof is administered in a dose of 10 to 50 µg per day.

17. (Previously presented) The method according to claim 16, wherein the dose is administered once daily.

18. -24. (Canceled)

25. (New) The pharmaceutical composition according to Claim 6, further comprising an excipient and/or vehicle.

26. (New) The pharmaceutical composition according to Claim 25, wherein the excipient and/or vehicle is lactose.

27. (New) The pharmaceutical composition according to Claim 25, wherein the excipient and/or vehicle is lactose monohydrate.

28. (New) The method according to Claim 11, wherein the pharmaceutical composition further comprises an excipient and/or vehicle.

29. (New) The method according to Claim 28, wherein the excipient and/or vehicle is lactose.
30. (New) The method according to Claim 28, wherein the excipient and/or vehicle is lactose monohydrate.